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(54) **Magnesium salt of s-omeprazole**

(57) A process of producing the magnesium salt of an enantiomer of omeprazole, said process comprising the steps of:

vii) reacting magnesium with a lower alcohol to produce magnesium alkoxide in solution in the lower alcohol as solvent,

viii) adding the neutral form of the enantiomer of omeprazole to the solution, and
ix) flash-evaporating the solvent.

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Description

[0001] The present invention relates to an improved form of the magnesium salt of S-omeprazole, a process for making same, and pharmaceutical compositions thereof.

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BACKGROUND OF THE INVENTION

[0002] The compound known under the generic name omeprazole is described in European patent 0005129. Further, European Patent No. 124,495 describes the advantages of providing the salts of omeprazole and particularly the magnesium salt thereof.

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[0003] Omeprazole is useful for inhibiting gastric acid secretion and has gastric mucosa protective activity in mammals and man. Omeprazole may be used for prevention and treatment of gastric acid related disorders and gastrointestinal inflammatory diseases in mammals and man, including for example gastritis, gastric ulcer and duodenal ulcer.

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[0004] The terms "omeprazole, S-omeprazole and R-omeprazole" as used in this specification designate the neutral form thereof, that is the form without a salt-forming cation present, unless otherwise indicated.

[0005] European Patent No. 0124495, in example 5 specifically discloses the synthesis of magnesium omeprazole dihydrate, and example 6 specifically discloses the synthesis of magnesium omeprazole anhydrate. Manufacturing of the described magnesium omeprazole salt presents significant difficulties.

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[0006] The process of manufacture and isolation of the dihydrate according to example 5 is relatively complex. It requires making the sodium salt, adding a solution of magnesium chloride to obtain a precipitate, removing water by centrifuging the precipitate, washing the precipitate with deionized water until no Cl⁻ is detectable, drying in air, grinding, and then drying in vacuum at 40°C for 24h. Moreover, because the resulting magnesium omeprazole dihydrate is crystalline, the rate of dissolution in intestinal fluid is relatively slow, unless the material is milled to a relatively fine particle size. It would therefore be desirable to provide non-crystalline forms to improve the dissolution in intestinal fluid.

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[0007] The process of making the anhydrate according to example 6 is simpler. Magnesium is reacted with methanol to give a solution of magnesium methoxide in methanol. The solution is added to a solution of omeprazole in methanol, the quantity of omeprazole being one mole for each two moles of magnesium. The methanol is then evaporated to give a crystalline solid, which is magnesium omeprazole anhydrate. However, the anhydrate as made by this process is also not without a problem. As the magnesium omeprazole precipitates from the solution upon evaporation of the methanol, residual methanol is entrapped in the solid particles and cannot easily be removed by evaporation. Methanol is toxic and high levels are generally considered unacceptable in pharmaceutical chemicals.

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[0008] Canadian patent 2166794 describes what is said to be an improved form of magnesium omeprazole dihydrate, which has a higher degree of crystallinity than that of example 5 of EP 0124495. This form has a methanol content of less than 0.1%. However, like the product of example 6 of EP 0124495, it is a crystalline dihydrate, and the process of manufacture is relatively complex.

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[0009] According to Canadian patent 2166794, the degree of crystallinity of a sample made according to example 6 of EP 0124495 was 67%, whereas the degree of crystallinity of the improved form is at least 70%.

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[0010] Canadian patent application No. 2254572 discloses improved processes for the production of magnesium omeprazole crystalline dihydrate. The disclosure reviews the prior art, and in particular, in relation to the anhydrate of example 6 of EP 0124495, states as follows: *"This procedure cannot be practiced on a large scale because of the need to evaporate to dryness. It has been found that unacceptable and potentially dangerous amounts of methanol become trapped in this solid, making it pharmaceutically unacceptable."* The processes of Canadian patent 2254572 are again relatively complex.

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[0011] Improved processes for the production of magnesium omeprazole crystalline dihydrate are also described in PCT Publication No. WO 97/41114. The degree of crystallinity of the product of example 1 is said to be 80%. Again, the processes disclosed are relatively complex.

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[0012] Omeprazole is a sulfoxide and a chiral compound, wherein the sulfur atom is the stereogenic center. Thus, omeprazole is a racemic mixture of its two single enantiomers, the R and S-enantiomer of omeprazole, herein referred to as R-omeprazole and S-omeprazole. The absolute configurations of the enantiomers of omeprazole have been determined by an X-ray study of an N-alkylated derivative of the (+)-enantiomer in non-salt form. The (+)-enantiomer of the non-salt form and the (-)-enantiomer of the non-salt form were found to have R and S configuration, respectively, and the (+)-enantiomer of the magnesium salt and the (-)-enantiomer of the magnesium salt were also found to have R and S configuration, respectively. The conditions for the optical rotation measurement for each of these enantiomers are described in WO 94/27988.

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[0013] Certain salts of single enantiomers of omeprazole and their preparation are disclosed in WO 94/27988. These compounds have improved pharmacokinetic and metabolic properties which will give an improved therapeutic profile such as a lower degree of interindividual variation.

[0014] WO 96/02535 discloses a process for the preparation of the single enantiomers of omeprazole and salts

thereof, and WO 96/01623 discloses a suitable tableted dosage form for instance magnesium salts of R- and S-omeprazole. The magnesium salt of S-omeprazole trihydrate described is substantially free from magnesium salts of R-omeprazole.

5 [0015] United States Patent No. 5,714,504 describes optically pure salts of omeprazole and in particular the sodium and magnesium salts thereof as pure crystalline enantiomeric salts, and in one embodiment optically pure crystalline magnesium salts. The patent describes the non-aqueous process for the preparation of crystalline forms of the magnesium salts of optically pure enantiomers of omeprazole or analogues thereof; which include the steps of stirring a crude preparation of the omeprazole enantiomer under nitrogen into a methanolic magnesium methoxide solution, precipitating inorganic magnesium salts with the addition of a small amount of water, removing any precipitated inorganic magnesium salts, concentrating the residual methanolic solution, precipitating the omeprazole enantiomer by adding acetone to the residual solution, and filtering off the optically pure enantiomer crystals of magnesium omeprazole or analogues thereof. Because it is possible to purify optically impure or partially pure salts of the enantiomers of omeprazole by crystallization, they can be obtained in very high optically pure, namely greater than or equal to 99.8% enantiomeric excess. Example 6 within the specification describes the preparation of the magnesium salt of S-omeprazole by crystallization of said salt.

10 [0016] The preferred enantiomer of omeprazole referred to as the (-)-enantiomer of omeprazole or a pharmaceutical salt thereof, is said to be an improved alternative to omeprazole in the treatment of gastric acid related diseases which provides higher dose efficiencies and less inter-individual variation in plasma levels, both between rapid and slow metabolizers and within the group of rapid metabolizers, as taught in United States Patent No. 5,877,192. The major 15 emphasis described relates to various forms of the enantiomers of omeprazole and salts thereof in crystalline form and preferably in highly crystalline form, which are also described in Canadian Patent Application No. 2,357,744. Although amorphous forms are nominally discussed there is no specific teaching as to the advantages of preventing crystals from forming. Therefore, a need exists for the magnesium salts of enantiomers of omeprazole having a desirable low methanol content.

20 [0017] United States Patent No. 6,262,085 teaches in example 20 the magnesium salt of S-omeprazole. Generally, the patent describes the preferred crystalline form but states that other forms such as amorphous forms are casually mentioned, but clearly the teaching refers to crystalline and particularly to the co-crystalline form wherein enantiomers of omeprazole are present in the same crystal lattice and co-crystallized from solution. However, there is no teaching 25 as to the manner in which amorphous forms in particular might be prepared, resulting in the same deficiencies with reference to solvent content as described above.

30 [0018] It would therefore be highly desirable to provide primarily amorphous magnesium salt of the enantiomers of omeprazole and particularly the magnesium salt of S-omeprazole, since these salts have surprisingly high stability in alkaline conditions. There still exists a need for magnesium salts of enantiomers of omeprazole having substantially 35 low methanol content and having a minimum amount of crystallinity with a large percentage of the material being amorphous, that is having minimum crystalline structure.

[0019] In summary, the only magnesium omeprazole according to the prior art that has an acceptably low level of methanol is magnesium omeprazole crystalline dihydrate, which has a degree of crystallinity of 67% or higher and is produced only by relatively complex processes.

40 [0020] In light of the foregoing, the object of the present invention is to produce magnesium omeprazole and the magnesium salt of enantiomers of omeprazole having acceptably low levels of methanol, but containing a large proportion of amorphous material (non-crystalline), which preferably may also be substantially amorphous as well, to be produced by a simple process.

45 [0021] It is also an object of this invention to provide the magnesium salt of S-omeprazole in pharmaceutically acceptable forms.

[0022] It is a further object of the invention to provide the magnesium salt of R-omeprazole in pharmaceutically acceptable forms.

50 [0023] Further and other objects of the invention will become apparent to those skilled in the art when considering the following summary of the invention and the more detailed description of the preferred embodiments and examples contained herein.

BRIEF SUMMARY OF THE INVENTION

55 [0024] According to one aspect of the invention magnesium omeprazole of the present invention is made by reacting magnesium in a lower alcohol to form magnesium alkoxide, preferably adding omeprazole in a quantity of about two moles per mole of magnesium, and flash-evaporating the alcohol, so as to form a solid precipitate without allowing the growth of crystals or particles that entrap the alcohol at unacceptable levels. The resulting material is substantially amorphous (non-crystalline).

[0025] According to a primary aspect of the invention the magnesium salt of the enantiomers of omeprazole of the

present invention is made by reacting magnesium in a lower alcohol to form magnesium alkoxide, adding only one of the enantiomers of omeprazole in neutral form, for example S-omeprazole, or alternatively R-omeprazole, preferably in a quantity of about 2 moles per mole of magnesium, and flash evaporating the alcohol, so as to form a solid precipitate without allowing the substantial growth of crystals or particles that entrap the alcohol at unacceptable levels. The resulting material contains a desirable level of non-crystalline material, and preferably a primarily amorphous amount of the magnesium salts of either of the enantiomers of omeprazole, and preferably magnesium S-omeprazole. In another embodiment a substantially amorphous form is provided.

[0026] According to yet another aspect of the invention there is provided a process of producing the magnesium salt of an enantiomer of omeprazole, said process comprising the steps of:

- 10 i) reacting magnesium with a lower alcohol to produce magnesium alkoxide in solution in the lower alcohol as solvent,
- ii) adding the neutral form of the enantiomer of omeprazole to the solution, and
- iii) flash-evaporating the solvent.

[0027] In one embodiment the enantiomer is S-omeprazole. In another embodiment the enantiomer is R-omeprazole. Preferably the lower alcohol is methanol.

In one embodiment the flash-evaporation is done by spray-drying the solution.

[0028] According to yet another aspect of the invention there is provided magnesium S-omeprazole or alternatively magnesium R-omeprazole having a residual organic solvent content of less than 7% by weight.

[0029] According to yet another aspect of the invention there is provided magnesium S-omeprazole or alternatively magnesium R-omeprazole having a degree of crystallinity of under 67% and in one embodiment having a residual organic solvent content of less than 7% by weight, preferably a residual organic solvent content of less than 5% by weight, more preferably a residual organic solvent content of less than 2% by weight, and most preferably a residual organic solvent content of less than 1% by weight.

[0030] In one embodiment magnesium S-omeprazole or alternatively magnesium R-omeprazole has a degree of crystallinity of under 60%, preferably has a degree of crystallinity of under 50%, more preferably has a degree of crystallinity of under 25%.

[0031] Preferably a solid pharmaceutical composition for oral administration may further comprise magnesium S-omeprazole or alternatively magnesium R-omeprazole as described above, preferably in the form of a tablet, wherein the tablet may be enteric coated. In one embodiment the enteric coated tablet may further comprise a separating layer between said enteric coating and said tablet.

[0032] The resulting composition comprising magnesium S-omeprazole or alternatively magnesium R-omeprazole is preferably in substantially amorphous form.

DETAILED DESCRIPTION OF THE INVENTION

[0033] In the process of manufacture of magnesium omeprazole, or the magnesium salt of s-omeprazole according to one aspect of the present invention, magnesium is reacted in a lower alcohol, preferably methanol, to form a solution of magnesium alkoxide in the alcohol.

[0034] The atomic weight of magnesium is 24.3 and the molecular weight of omeprazole or the neutral form of s-omeprazole is 345.4. Since magnesium is divalent, the amount of magnesium required to convert 345.4 grams of omeprazole or S-omeprazole to magnesium omeprazole or the magnesium salt of s-omeprazole is 12.15 grams.

[0035] Hence 35.2 grams of magnesium is needed to convert 1 kilo of omeprazole or the neutral form of s-omeprazole to magnesium omeprazole or magnesium s-omeprazole.

[0036] The process of converting 1 kilo of omeprazole or the neutral form of s-omeprazole to magnesium omeprazole or the magnesium salt of s-omeprazole, thus begins with reacting 35.2 grams of magnesium in a lower alcohol, preferably methanol. The minimum amount of methanol needed to react fully and dissolve 35.2 grams of magnesium is about 1000 grams.

[0037] When the magnesium is immersed in the alcohol, the reaction will be evident from the generation of hydrogen bubbles, and the reaction will be complete when all the magnesium has been consumed and the effervescence has ceased. All of the magnesium will then be present as magnesium alkoxide in the alcohol (i.e. magnesium methoxide in methanol, if methanol is used as the alcohol).

[0038] The omeprazole or the neutral form of s-omeprazole can then be added directly to the magnesium alkoxide solution. Alternatively, the omeprazole or the s-omeprazole (neutral form) may first be dissolved in an alcohol or another organic solvent that is miscible with the alcohol used to make the magnesium alkoxide, and the resultant solution may then be added to the magnesium alkoxide solution.

[0039] Where methanol is used as the sole solvent, a total of only about 1.5 kilos is needed for converting 1 kilo of

omeprazole or the neutral form of S-omeprazole to magnesium omeprazole or the magnesium salt of S-omeprazole. [0040] Hence, using quantities based on 1 kilo of omeprazole or S-omeprazole (or alternatively R-omeprazole), the simplest and best procedure is to react 35.2 grams of magnesium in about 1.5 kilos of methanol, wait until the magnesium has been fully reacted, and then adding 1 kilo of omeprazole or S-omeprazole to the solution and stir to dissolve. 5 The result will be a solution of magnesium omeprazole or S-omeprazole equivalent to 1 kilo of omeprazole in methanol. [0041] In order to obtain solid, magnesium omeprazole or magnesium S-omeprazole that is substantially free of organic solvent (i.e. substantially free of methanol, if methanol is used), it is then necessary to eliminate the solvent. [0042] It has been found that this can be done according to one aspect of the invention by "flash-evaporating" the solvent. Flash-evaporating will be understood to mean evaporating in such a way as to avoid the precipitation of crystals or large particles which entrap the alcohol. 10 [0043] One method of flash-evaporating the solvent is to mix the solution into a solid excipient such as, for example, microcrystalline cellulose or the like, or any other well known appropriate excipient, so that a damp mass is formed. The mass can then be dried in a conventional oven, a fluid bed drier, or under vacuum to remove the solvent. Because the solution has been dispersed throughout the solid excipient, as the solvent evaporates, the magnesium omeprazole or the magnesium salt of S-omeprazole, is deposited as a thin layer over the surface of the particles of the solid excipient and does not precipitate as crystals or large granules, so that there is little or no entrapment of solvent. 15 [0044] The preferred way of flash-evaporating the solvent is by spray-drying the solution. [0045] It has been found in utilizing the above-mentioned preferred processes that magnesium omeprazole and the magnesium salt of S-omeprazole can be made having a residual solvent content substantially lower than can be 20 achieved by simply evaporating the solvent from the solution under vacuum. [0046] The residual organic solvent content by weight of the magnesium omeprazole, and the magnesium salt of S-omeprazole made according to the present invention will be under 7%, preferably under 5%, more preferably under 2%, and most preferably under 1%. [0047] The degree of crystallinity of the obtained product can be measured with powder X-ray diffraction (XRD) as 25 described in WO 97/41114 as follows: A thin layer of the triturated sample is smeared onto a cut silicon single crystal zero background holder which is rotated during the measurement. Cu KV radiation and constant or automatic antiscatter and divergence slits are used to obtain a diffractogram with 2θ from 1 or 2° to at least 35° . [0048] The degree of crystallinity is calculated with the formula: 30

$$\text{Degree of crystallinity} = 100 \cdot C / (A+C)$$

C= the area from the peaks in the diffractogram ("the crystalline area"),
 A= the area between the peaks and the background ("the amorphous area").

35 [0049] Area calculations are performed for 2θ between $4-33^\circ$. The lowest intensity value found in this interval is chosen as the constant background and subtracted from the area A. When constant slits are used, the increased background at low angles due to the influence from the primary beam is also subtracted from the area A. [0050] The degree of crystallinity of magnesium omeprazole and the magnesium salt of S-omeprazole according to 40 the present invention is under 67%, as compared to 67% or higher for magnesium omeprazole crystalline dihydrate according to the prior art. [0051] The degree of crystallinity will preferably be under 60%, more preferably under 50%, and most preferably under 25%. [0052] If the magnesium omeprazole or the magnesium salt of S-omeprazole of the present invention is made in an environment and using excipients (including the air or other gas used for drying in the spray-dry process) that is completely free of water, the magnesium omeprazole or the magnesium S-omeprazole will be anhydrous. However, pure anhydrous magnesium omeprazole or magnesium S-omeprazole is hygroscopic and it will readily absorb water from air until it reaches an equilibrium water content of about 5% to 8%, depending on the relative humidity of the air. This is not problematic, as it does not adversely affect stability of the final product. 45 [0053] The present invention will be further processed into pharmaceutical compositions such as, for example, tablets for oral administration. The tablets will preferably be enteric coated to protect the magnesium omeprazole and magnesium S-omeprazole from the effects of gastric acid. [0054] The invention will be further understood from the following examples, which are intended to be illustrative and not limiting of the invention. 50

55 EXAMPLE 1

[0055] 1.76 g of pure magnesium was added to 800 g of methanol in a 1000 mL glass flask. The flask was closed with a loose-fitting stopper (loose to allow hydrogen gas to escape), and the flask was allowed to sit overnight.

5 [0056] The next morning it was observed that the magnesium had all been consumed and that the effervescence had ceased, resulting in a slightly hazy solution of magnesium methoxide in methanol. 50 grams of omeprazole (or the neutral form of S-omeprazole could be used) was then added to the contents of the flask and the contents were stirred for several minutes until dissolved to form a solution of magnesium omeprazole (or if the neutral form of S-omeprazole was used, magnesium S-omeprazole) in methanol.

EXAMPLE 2

10 [0057] To produce a reference sample of magnesium omeprazole anhydrate according to the prior art (i.e. example 6 of EP 0124495), about 20% of the solution from step 2 was transferred to a 1000 mL beaker. The beaker was then placed in a vacuum oven for drying under vacuum at 50°C for a period of 4 hours. At the end of this time, a solid material remained that had no evident odour of residual methanol. This solid material was tested to determine the level of residual methanol, which was found to be 7.2% by weight.

15 EXAMPLE 3

[0058] To produce the present invention, the balance of the solution of Example 1 was spray-dried on a Yamato® spray-dryer, using an inlet air temperature of about 140°C and outlet air temperature of about 70°C.

20 [0059] The resulting dry material was a fine powder, which appeared non-crystalline (i.e. amorphous) and also had no evident odour of residual methanol. The powder was tested to determine the level of residual methanol, which was found to be 0.7%.

[0060] This powder was examined for crystallinity by powder X-ray diffraction, and it was found that the powder was primarily amorphous (non-crystalline), having a degree of crystallinity of under 25%.

25 EXAMPLE 4

[0061] The following ingredients are to be mixed together in the proportions shown:

30	Magnesium S-omeprazole (prepared according to Examples 1 and 3)	21.0
	Anhydrous lactose	131.0
35	Croscarmellose sodium	6.4
	Magnesium stearate	1.6
		<hr/> 160.0

40 [0062] The mixture is to be compressed into tablets having a weight of 160 mg per tablet, so that each tablet will contain 21 mg of magnesium S-omeprazole, which is equivalent to about 20 mg of omeprazole.

[0063] A sub-coating comprising hydroxypropyl methylcellulose dissolved in water will then be applied to the tablets by spray-application in a side-vented coating pan.

45 [0064] An enteric coating is to be applied over the sub-coating by spray-application of methacrylic acid copolymer aqueous dispersion, with triethyl citrate dissolved therein as plasticizer.

[0065] As many changes can be made to the invention without departing from the scope of the invention, it is intended that all material contained herein be interpreted as illustrative of the invention and not in a limiting sense.

50 **Claims**

1. A process of producing the magnesium salt of an enantiomer of omeprazole, said process comprising the steps of:

55 iv) reacting magnesium with a lower alcohol to produce magnesium alkoxide in solution in the lower alcohol as solvent,
v) adding the neutral form of the enantiomer of omeprazole to the solution, and
vi) flash-evaporating the solvent.

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2. The process of claim 1 wherein the enantiomer is S-omeprazole.
3. The process of claim 1 wherein the enantiomer is R-omeprazole.
5. The process of claim 1, 2 or 3 wherein the lower alcohol is methanol.
10. The process of claim 1, 2, 3 or 4 wherein the flash-evaporation is done by spray-drying the solution.
6. Magnesium S-omeprazole or magnesium R-omeprazole having a residual organic solvent content of less than 7% by weight.
15. Magnesium S-omeprazole or magnesium R-omeprazole having a degree of crystallinity of under 67%.
8. Magnesium S-omeprazole or magnesium R-omeprazole of claim 7 having a residual organic solvent content of less than 7% by weight.
15. Magnesium S-omeprazole or magnesium R-omeprazole of claim 6 or 7 having a residual organic solvent content of less than 5% by weight.
20. Magnesium S-omeprazole or magnesium R-omeprazole of claim 6 or 7 having a residual organic solvent content of less than 2% by weight.
11. Magnesium S-omeprazole or magnesium R-omeprazole of claim 6 or 7 having a residual organic solvent content of less than 1% by weight.
25. Magnesium S-omeprazole or magnesium R-omeprazole of any of claims 6 to 11 having a degree of crystallinity of under 60%.
30. Magnesium S-omeprazole or magnesium R-omeprazole of any of claims 6 to 11 having a degree of crystallinity of under 50%.
14. Magnesium S-omeprazole or magnesium R-omeprazole of any of claims 6 to 11 having a degree of crystallinity of under 25%.
35. 15. A solid pharmaceutical composition for oral administration comprising magnesium S-omeprazole or magnesium R-omeprazole of any of claims 6 to 14.
16. The composition of claim 15 in the form of a tablet.
40. 17. The composition of claim 16 wherein the tablet is enteric coated.
18. The process or composition of claim 2, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16 or 17 wherein magnesium S-omeprazole or magnesium R-omeprazole is in substantially amorphous form.
45. 19. The composition of claim 17 further comprising a separating layer between said enteric coating and said tablet.



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EUROPEAN SEARCH REPORT

Application Number

EP 03 25 3070

DOCUMENTS CONSIDERED TO BE RELEVANT			CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	
X	WO 01 36409 A (SHERMAN BERNARD CHARLES) 25 May 2001 (2001-05-25)	1-19	C07D401/12 A61K31/44
Y	Claims 1 to 14 * the whole document * ---	1-19	//(C07D401/12, 25:00,213:00)
Y	WO 94 27988 A (ASTRA AB ; LINDBERG PER LENNART (SE); VON UNGE SVERKER (SE)) 8 December 1994 (1994-12-08) * page 4, line 19 - line 21; examples 3-5 * ---	1-19	
A,D	EP 0 124 495 A (HAESSLE AB) 7 November 1984 (1984-11-07) Page 2, pages 13 to 14 and table 1 ---	1	
A	WO 95 01977 A (ASTRA AB ; KAEILSTROEM LARS AAKE (SE); NYGREN MONICA ANNELIE (SE)) 19 January 1995 (1995-01-19) page 2, second paragraph -----	1	
The present search report has been drawn up for all claims			TECHNICAL FIELDS SEARCHED (Int.Cl.7)
			C07D A61K
Place of search		Date of completion of the search	Examiner
MUNICH		3 July 2003	Goss, I
CATEGORY OF CITED DOCUMENTS			
X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document			
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ON EUROPEAN PATENT APPLICATION NO.

EP 03 25 3070

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03-07-2003

Patent document cited in search report		Publication date		Patent family member(s)	Publication date
WO 0136409	A	25-05-2001	CA	2290893 A1	16-05-2001
			AU	6551100 A	30-05-2001
			WO	0136409 A1	25-05-2001
			EP	1230237 A1	14-08-2002
			JP	2003514813 T	22-04-2003
<hr/>					
WO 9427988	A	08-12-1994	AT	197452 T	11-11-2000
			AU	676337 B2	06-03-1997
			AU	6902494 A	20-12-1994
			CA	2139653 A1	08-12-1994
			CA	2337581 A1	08-12-1994
			CN	1110477 A ,B	18-10-1995
			CN	1259346 A ,B	12-07-2000
			CZ	9500202 A3	18-10-1995
			DE	69426254 D1	14-12-2000
			DE	69426254 T2	07-06-2001
			DE	652872 T1	04-09-1997
			DK	652872 T3	05-03-2001
			EE	3157 B1	15-02-1999
			EP	1029460 A2	19-07-2000
			EP	1020461 A2	19-07-2000
			EP	0652872 A1	17-05-1995
			ES	2099047 T1	16-05-1997
			FI	950377 A	27-01-1995
			GR	97300012 T1	31-05-1997
			GR	3035365 T3	31-05-2001
			HK	1008330 A1	06-07-2001
			HR	940307 A1	31-12-1996
			HU	71888 A2	28-02-1996
			IL	109684 A	23-05-2002
			JP	7509499 T	19-10-1995
			LT	1941 A ,B	27-12-1994
			LV	11034 A	20-02-1996
			LV	11034 B	20-10-1996
			NO	950263 A	24-01-1995
			NZ	266915 A	28-10-1996
			PL	307261 A1	15-05-1995
			PT	652872 T	30-04-2001
			RU	2137766 C1	20-09-1999
			WO	9427988 A1	08-12-1994
			SG	49283 A1	18-05-1998
			SI	9420002 A	31-08-1995
			SK	10195 A3	13-09-1995
			TW	389761 B	11-05-2000
			US	5693818 A	02-12-1997
			US	5714504 A	03-02-1998

EPO FORM P0459

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82

ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.

EP 03 25 3070

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report.
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03-07-2003

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9427988 A		US 6143771 A US 5877192 A ZA 9403557 A	07-11-2000 02-03-1999 11-04-1995
EP 0124495 A	07-11-1984	AT 24907 T AU 563842 B2 AU 2525784 A BG 44538 A3 BG 60837 B2 CA 1264751 A1 CS 8401515 A2 DD 221459 A5 DE 3462036 D1 DK 99584 A ,B, EP 0124495 A2 ES 8500934 A1 FI 840851 A ,B, GB 2137616 A ,B GR 79828 A1 HK 13590 A HR 930428 B1 HU 193557 B IE 57326 B1 IL 70985 A JP 1651336 C JP 3013233 B JP 59167587 A KR 8701005 B1 LT 2253 R3 LU 90677 A9 LV 5503 A3 LV 5801 A4 NO 840772 A ,B, NZ 207348 A PH 21352 A PL 246492 A1 PT 78191 A ,B RO 88721 A1 SG 1490 G SI 8410397 A8 SU 1314953 A3 US 4738974 A YU 39784 A1 ZA 8401202 A	15-01-1987 23-07-1987 06-09-1984 15-12-1988 30-04-1996 23-01-1990 13-06-1985 24-04-1985 19-02-1987 05-09-1984 07-11-1984 01-02-1985 05-09-1984 10-10-1984 31-10-1984 02-03-1990 30-04-1996 28-10-1987 29-07-1992 20-10-1987 30-03-1992 22-02-1991 21-09-1984 18-05-1987 15-11-1993 05-02-2001 10-03-1994 20-02-1997 05-09-1984 08-10-1986 15-10-1987 27-02-1985 01-04-1984 30-04-1986 13-07-1990 31-10-1995 30-05-1987 19-04-1988 31-12-1986 31-10-1984
WO 9501977 A	19-01-1995	AT 212628 T	15-02-2002

EU For more details about this annex : see Official Journal of the European Patent Office, No. 12/82

ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.

EP 03 25 3070

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

03-07-2003

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9501977 A			
	AU	679766 B2	10-07-1997
	AU	7198194 A	06-02-1995
	BR	9406940 A	10-09-1996
	CA	2166794 C	04-03-1997
	CN	1126993 A ,B	17-07-1996
	CZ	9600069 A3	15-05-1996
	DE	69429774 D1	14-03-2002
	DE	69429774 T2	14-11-2002
	DE	707580 T1	04-09-1997
	DK	707580 T3	15-04-2002
	EE	3127 B1	15-10-1998
	EG	21437 A	31-10-2001
	EP	1164132 A2	19-12-2001
	EP	0707580 A1	24-04-1996
	ES	2100136 T1	16-06-1997
	FI	960101 A	09-01-1996
	GR	97300015 T1	31-05-1997
	HK	1008329 A1	17-05-2002
	HR	940385 A1	28-02-1997
	HU	75314 A2	28-05-1997
	IL	110190 A	26-07-2000
	JP	8512315 T	24-12-1996
	JP	2002105072 A	10-04-2002
	MX	9405217 A1	31-01-1995
	NO	960068 A	05-01-1996
	NZ	268693 A	26-05-1997
	PL	312440 A1	29-04-1996
	PT	707580 T	28-06-2002
	RU	2139868 C1	20-10-1999
	WO	9501977 A1	19-01-1995
	SG	52464 A1	28-09-1998
	SI	707580 T1	30-06-2002
	SK	2296 A3	01-10-1996
	US	5900424 A	04-05-1999
	ZA	9404933 A	20-02-1995